FDA 510(k) Premarket Notification

K980213 DI-2000

CLEARED

Mar 17, 1998
67 days to decision

K980213 · Product code: **LMA** · Radiology
Source: https://www.510kdatabase.net/k980213/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digitizer, Image, Radiological (LMA)
Date received	Jan 9, 1998
Decision date	Mar 17, 1998
Days to decision	67 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lumisys, Inc.**
Location	San Leandro, CA, US
Contact	TRINDY LEFORGE
510(k) history	7 submissions · 7 cleared · 1990-1998